

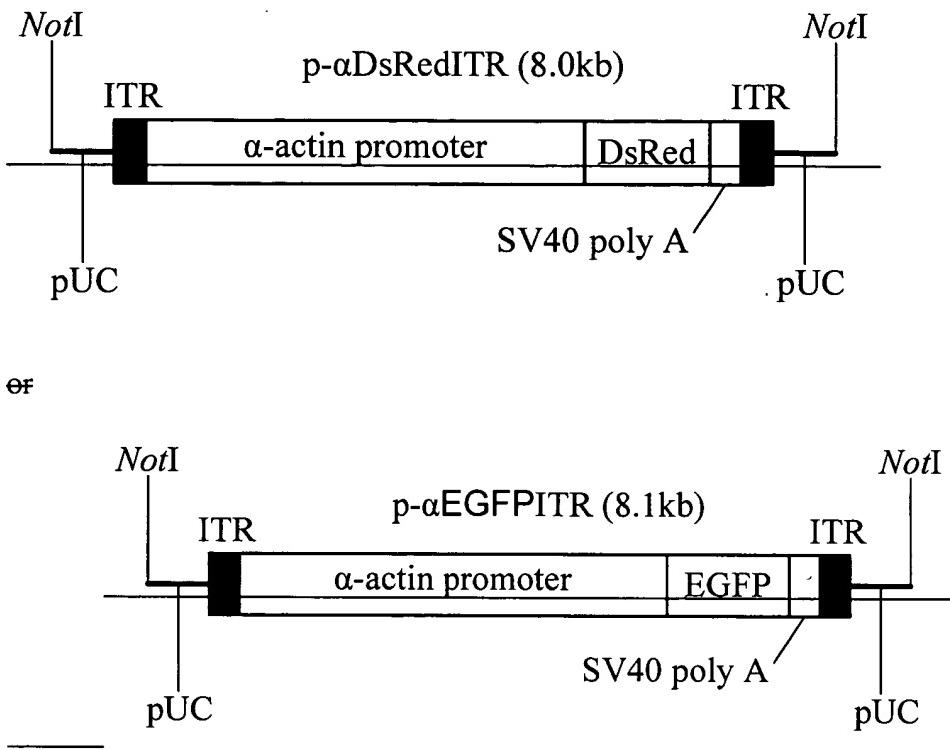
AMENDMENTS TO THE CLAIMS

The following is a complete, marked up listing of revised claims with a status identifier in parentheses, underlined text indicating insertions, and strikethrough and/or double-bracketed text indicating deletions.

LISTING OF CLAIMS

1. (CURRENTLY AMENDED) A gene fragment comprising, in an upstream to downstream order the operatively linked regions (1) first inverted terminal repeats (ITR) of adeno-associated virus; (2) an α -actin gene promoter of golden zebrafish; [(2)] (3) a gene encoding a red fluorescence gene product; (4) SV40 poly A and [(3)] (5) second inverted terminal repeats (ITR) of adeno-associated virus; and (4) ~~a basic part from pUC~~.

2. (CURRENTLY AMENDED) The gene fragment of Claim 1 further comprising which is—a first pUC backbone segment operatively linked to and upstream of the first inverted terminal repeats and a second pUC backbone segment operatively linked to and downstream of the second inverted terminal repeats and wherein the gene encoding the red fluorescence gene product is DsRed



3. (CURRENTLY AMENDED) A method of producing an adult golden zebrafish with systemic red fluorescence comprising:

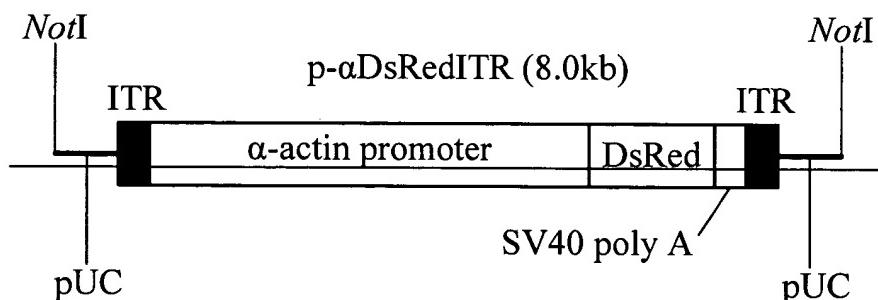
- (a) constructing a plasmid including a first ITR, a CMV promoter, a gene encoding a fluorescent gene product, SV40 poly A and a second ITR;
- (b) replacing the CMV promoter with an α -actin gene promoter of golden zebrafish to produce a new plasmid construct in which the α -actin gene promoter is operably linked to the gene encoding a fluorescent gene product;
- (c) linearizing the new plasmid construct;
- (d) microinjecting the linearized new plasmid construct into fertilized golden zebrafish eggs ~~of golden zebrafish~~ to obtain microinjected eggs;

(e) incubating the microinjected eggs for at least 24 hours to form embryos;

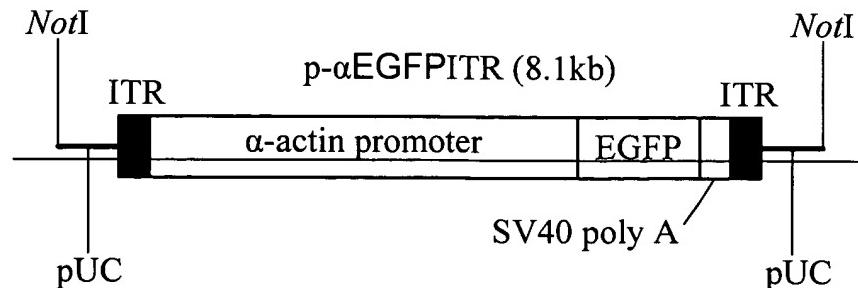
(f) selecting incubated an embryo eggs exhibiting red fluorescence; and

[(f)] (g) cultivating the selected embryo eggs to maturity to produce an adult golden zebrafish having skeletal muscle that exhibits red fluorescence.

4. (CURRENTLY AMENDED) The method of Claim 3 wherein the linearized plasmid is consists of, in upstream to downstream order, operably linked regions designated as a first pUC backbone segment, a first ITR, an α -actin gene promoter for golden zebrafish, a gene encoding a red fluorescent gene product, SV40 Poly A, a second ITR, and a second pUC backbone segment



or



5. (CURRENTLY AMENDED) The method of Claim 3 wherein the gene encoding the red fluorescent gene product is a red fluorescent gene from pDsRed2-1.

6. (CANCELED)

7. (CURRENTLY AMENDED) [[A]] An adult golden zebrafish having skeletal muscle that exhibits systemic red fluorescence produced according to the method of Claim 3.

8. (CURRENTLY AMENDED) The adult golden zebrafish of Claim 7 wherein the linearized plasmid consists of, in upstream to downstream order, operably linked regions designated as a first pUC backbone segment, a first ITR, an α-actin gene promoter for golden zebrafish, a gene encoding a red fluorescent gene product, SV40 Poly A, a second ITR, and a second pUC backbone segment in which skeletal muscle exhibits red fluorescence.

9. (CANCELED)

10. (CURRENTLY AMENDED) The method of Claim 3, wherein the linearized plasmid is selected from a group consisting of
a first-linearized plasmid consisting of, in order, a first pUC backbone segment, a first ITR, an α-actin gene promoter for of golden zebrafish, a gene encoding a red fluorescent gene product, SV40 Poly A, a second ITR, and a second pUC backbone

wherein the first ITR is located at a 5' end of the α -actin gene promoter and the second ITR is located at a 3' end of the SV40 poly A, further wherein the gene encoding a red fluorescent gene product and the gene promoter are operably linked, and further wherein the first and second pUC backbone segments may be cut with *NotI*;

and

~~a second linearized plasmid consisting of, in order, a first pUC backbone segment, a first ITR, an α -actin gene promoter for golden zebrafish, gene encoding a green fluorescent gene product, SV40 Poly A, a second ITR, and a second pUC backbone segment, wherein the first ITR is located at a 5' end of α -actin gene promoter and the second ITR is located at a 3' end of the SV40 poly A, wherein the gene encoding a green fluorescent gene product and the gene promoter are operably linked, and further wherein the first and second pUC backbone segments may be cut with *NotI*.~~

11. (CURRENTLY AMENDED) The method of Claim 10, wherein:

— the gene encoding a red fluorescent gene product is DsRed; and
~~gene encoding a green fluorescent gene product is EGFP.~~

12. (CURRENTLY AMENDED) The method of Claim 3 wherein the linearized plasmid is selected from a group consisting of

 a first linearized plasmid consisting of, in order, a first pUC backbone segment, a first ITR, an α -actin gene promoter capable of activity in ~~for~~ golden zebrafish, a ~~the~~ gene encoding a red fluorescent gene product, SV40 Poly A, a second ITR, and a second pUC backbone wherein the first ITR is located at a 5' end of the α -actin gene promoter and the

second ITR is located at a 3' end of the SV40 poly A, wherein the gene encoding a red fluorescent gene product and the gene promoter are operably linked, and further wherein the first and second pUC backbone segments may be cut with *NotI*.

13. (CANCELED)

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